

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

DMB

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21 CFR Part 176

[Docket No. 99F-1719]

Indirect Food Additives: Paper and Paperboard Components

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the food additive regulations to provide for the safe use of 4-(diiodomethylsulfonyl) toluene as a slimicide in the manufacture of food-contact paper and paperboard. This action is in response to a petition filed by Angus Chemical Co.

DATES: This rule is effective [*insert date of publication in the Federal Register*]. Submit written objections and requests for a hearing by [*insert date 30 days after date of publication in the Federal Register*].

ADDRESSES: Submit written objections to the Dockets Management Branch (HFA-305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Mark A. Hepp, Center for Food Safety and Applied Nutrition (HFS-215), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202-418-3098.

SUPPLEMENTARY INFORMATION: In a notice published in the **Federal Register** of June 11, 1999 (64 FR 31593), FDA announced that a food additive petition (FAP 9B4668) had been filed by Angus Chemical Co., c/o Phillip A. Johns, 10900 Silent Wood Pl., North Potomac, MD 20878-4829. The petition proposed to amend the food additive regulations in § 176.300 *Slimicides* (21

CFR 176.300) to provide for the safe use of 4-(diiodomethylsulfonyl) toluene as a slimicide in the manufacture of food-contact paper and paperboard.

FDA has evaluated data in the petition and other relevant material. Based on this information, the agency concludes that: (1) The proposed use of the additive as a slimicide in the manufacture of food-contact paper and paperboard is safe, (2) the additive will achieve its intended technical effect, and therefore, (3) the regulations in § 176.300 should be amended as set forth below.

In accordance with § 171.1(h) (21 CFR 171.1(h)), the petition and the documents that FDA considered and relied upon in reaching its decision to approve the petition are available for inspection at the Center for Food Safety and Applied Nutrition by appointment with the information contact person listed above. As provided in § 171.1(h), the agency will delete from the documents any materials that are not available for public disclosure before making the documents available for inspection.

The agency has carefully considered the potential environmental effects of this rule as announced in the notice of filing for the petition. No new information or comments have been received that would affect the agency's previous determination that there is no significant impact on the human environment and that an environmental impact statement is not required.

This final rule contains no collection of information. Therefore, clearance by the Office of Management and Budget under the Paperwork Reduction Act of 1995 is not required.

Any person who will be adversely affected by this regulation may at any time file with the Dockets Management Branch (address above) written objections by *[insert date 30 days after date of publication in the **Federal Register**]*. Each objection shall be separately numbered, and each numbered objection shall specify with particularity the provisions of the regulation to which objection is made and the grounds for the objection. Each numbered objection on which a hearing is requested shall specifically so state. Failure to request a hearing for any particular objection shall constitute a waiver of the right to a hearing on that objection. Each numbered objection for which a hearing is requested shall include a detailed description and analysis of the specific

factual information intended to be presented in support of the objection in the event that a hearing is held. Failure to include such a description and analysis for any particular objection shall constitute a waiver of the right to a hearing on the objection. Three copies of all documents are to be submitted and are to be identified with the docket number found in brackets in the heading of this document. Any objections received in response to the regulation may be seen in the Dockets Management Branch between 9 a.m. and 4 p.m., Monday through Friday.

List of Subjects in 21 CFR Part 176

Food additives, Food packaging.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, and redelegated to the Director, Center for Food Safety and Applied Nutrition, 21 CFR part 176 is amended as follows:

PART 176—INDIRECT FOOD ADDITIVES: PAPER AND PAPERBOARD COMPONENTS

1. The authority citation for 21 CFR part 176 continues to read as follows:

Authority: 21 U.S.C. 321, 342, 346, 348, 379e.

2. Section 176.300 is amended in the table in paragraph (c) by alphabetically adding an entry under the headings "List of substances" and "Limitations" to read as follows:

§ 176.300 Slimicides.

* * * * *

(c) * * *

List of substances	Limitations
4-(Diiodomethylsulfonyl) toluene (CAS Reg. No. 20018-09-01).	At a maximum level of 0.2 pound per ton (100 grams/1,000 kilograms) of dry weight fiber.

* * * * *

Dated: 11/14/00
November 14, 2000

cf0090



L. Robert Lake
Director of Regulations and Policy
Center for Food Safety and
Applied Nutrition

CERTIFIED TO BE A TRUE
COPY OF THE ORIGINAL



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